Certificate of Conformity

VWR Catalogue Number	See List
Description	Disposable High Performance Transfer pipettes
Country of Origin	Manufactured in Italy
Date of Issue (yyyy-mm-dd)	2025-07-29

General Purpose

Catalogue Number	Capacity [mL]	Stem Diameter [µl]	Length [mm]	Sterile
612-2846	1.5	5	87	Non-Sterile
612-2849	1.5	5	123	Non-Sterile

Graduated

Catalogue Number	Capacity [mL]	Stem Diameter [µl]	Length [mm]	Graduation [mL]	Sterile
612-2851	3.0	5	138	1.0	Non-Sterile
612-1684	5.0	5	150	1.0	Non-Sterile
612-1685	5.0	5	150	1.0	Sterile
612-1686	5.0	5	150	1.0	Sterile
612-1687	5.0	5	150	1.0	Sterile
612-2842	6.0	5	230	1.5	Non-Sterile
612-2843	6.0	5	230	1.5	Sterile
612-2844	6.0	5	230	1.5	Sterile
612-2845	6.0	5	230	1.5	Sterile
612-1747	7.0	7.8	150	3.0	Sterile
612-1681	7.0	7.8	150	3.0	Non-Sterile
612-1683	7.0	7.8	150	3.0	Sterile
612-1682	7.0	7.8	150	3.0	Sterile

Extra Large

Catalogue Number	Capacity [mL]	Stem Diameter [µl]	Length [mm]	Sterile
612-2847	14.0	4	170	Non-Sterile
612-2859	23.0	9	300	Non-Sterile

Narrow Stem

Catalogue Number	Capacity [mL]	Stem Diameter [µl]	Length [mm]	Sterile
612-2848	1.2	2.5	63	Non-Sterile
612-2841	4.0	2.5	83	Non-Sterile
612-1688	4.0	2.5	150	Non-Sterile
612-1756	4.0	2.5	150	Sterile
612-1689	4.0	2.5	150	Sterile
612-1690	4.0	2.5	150	Sterile

Blood Bank

Catalogue Number	Capacity [mL]	Stem Diameter [µl]	Length [mm]	Graduation [mL]	Sterile
612-2850	5.0	6.7	155	2.0	Non-Sterile

Quality System Compliance

This certificate confirms that the product has been sourced from a supplier or manufacturer who has declared their Quality Management System (QMS) complies with **ISO 13485:2016.**

QC Testing

Representative products samples are collected and inspected in accordance with current applicable QC requirements.

Product Specifications

Material	Polyethylene. The materials used to produce the products are listed in List of Authorised Substances of Regulation EU/10/2011. Commission Regulation (EU) No. 10/2011 as amended. The material grades conform to EU Directive 1935/2004 and 2023/2006 on materials and articles intended to come into contact with food.
Sterilization	Product labelled as sterile is EO treated and they meet a minimum requirement of 10^{-6} Sterility assurance level. Sterilisation procedure is validated in conformity with ISO 11135:2014 which is not applicable for Non-sterile products.
ATP Assay	Not applicable.
DNase & RNase Free	Not applicable.
BSE/TSE	No use of any raw material produced from or substances derived from animal origin. The manufacturing process of the product does not use any ingredient of animal origin and no material derived from or exposed to animals affected by or under quarantine for transmitting Animal Bovine Spongiform Encephalopathy/Spongiform Encephalopathy (BSE)/(TSE).
Cytotoxicity	Not applicable.
Non-Pyrogenic Statement	Not applicable.
Latex Statement	This product is not made from natural rubber latex.
BPA Statement	Not applicable.
DEHP Statement	Not applicable.
RoHS	Not applicable.
REACH Statement	No substances as listed on >Substance of Very High Concern< candidate list published 2022-01-17 are used in manufacturing of the raw materials used to produce this product. No routinely analyse is performed.
Storage Conditions	No special storage condition required.

Shelf Life	Products labelled as sterile has shelf life of 36 months whereas non – sterile products has shelf life of 60 months.

Disclaimer: VWR's Quality Management System (QMS) is ISO 9001:2015 certified as a wholesale distributor of finished products, ensuring effective oversight of finished product handling, traceability, and distribution processes. VWR is not a manufacture and does not engage in manufacturing, co-manufacturing or processing activities such as design, packaging, labeling, testing or quality release. Consequently, the supplier or manufacturer's QMS certification may differ from VWR's.

VWR states that this declaration will not discharge the user from their obligation to ensure the product is suitable for the intended use. This product is intended for research use only. It is not for use in diagnostic procedures, therapeutic applications, or any human or clinical use.

This document has been produced electronically and is valid without a signature.

Version History

Sl.no	Version number	Description	Month & Year
1	1.0	Created General CoQ using new layout	04-2021
2	1.1	Modified FDA information	04-2021
3	1.1	Internal Change-Deleted product ranges-612-2853, 612-2854, 612-2855, 612-4464, 612-4465, 612-2856, 612-2857, 612-2858, 612-2852	09-2024
2	2.0	Modified Quality system compliance and disclaimer	07-2025